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(54) Synergistic polyamide stabilization method

(57) Disclosed is a process for stabilizing polyamides against the damage effected by light, heat and/or oxidation while improving the differential dyeability, and processing stability therefore, comprising incorporating therein by melt-processing a synergistic additive system resulting in a uniform and intimate mixture with said polyamide, said additive system is selected from the group consisting of (I), (II) and (III)

Also disclosed are polyamide articles selected from the group consisting of molded articles, extruded articles and fibers, which optionally contain organic dyes, exhibiting outstanding color brightness and color stability.

(I) is from 0.01% to 1% by weight of a sterically hindered phenyl phosphonite (Ia) with from 0.05% to 5% by weight of oxanilide (Ib) or said (Ia) together with from 0.05% to 5% by weight of an aromatic di- or tri-carbonyl compound containing at least one, and preferably two hindered amine moieties (Ic);

(II) is from 0.05% to 5% by weight of oxanilide (IIa) together with from 0.05% to 5% by weight of a aromatic di- or tri-carbonyl compound (IIb) containing at least one hindered amine moieties;

(III) is from 0.01% to 1% by weight of a sterically hindered phenyl phosphonite (Ia), from 0.05% to 5% by weight of a oxanilide (Ib) and from 0.05% to 5% by weight of an aromatic di- or tri-carbonyl compound;

wherein the percentages represent grams of additives used per hundred grams of polymer.

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Description**FIELD OF THE INVENTION**

- 5 [0001] The invention pertains to methods to stabilize polyamide polymers.

BACKGROUND OF INVENTION

[0002] The service life of polyamides is dependent on (1) the raw materials used in the manufacturing process, (2) the additives used in stabilizing the polymer against damage brought about by melt-processing and (3) environmental weathering factors. A serious drawback to providing stabilization systems for polyamides lies in the fact that the maximum concentration of many additives is limited. Polyamides are heretofore regarded as poor in solvating additives, especially non-polar additives having relatively low molecular weight in relation to the polyamide polymer. The aesthetics of an article such as made by molding processes can be affected such as by migration, blooming or plate-out of the additives. On the other hand, manufacturers of polyamide-based molded and extruded goods strive to include low concentrations of additives for economic reasons.

[0003] Oxalic acid di-aryl anilides, formed on condensing the acid, partial ester or di-ester with one or two moles of aniline derivatives are known light stabilizers. Unsymmetrical oxalic acid dianilides are described, for example in U.S. Pat. No. 4,003,875.

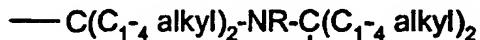
20 [0004] U.S. Pat. No. 5,045,083 discloses polyamide, copper complex and oxanilide UV absorber.

[0005] U.S. Pat. No. 4,544,691 discloses stabilizer compounds containing UV-absorbing derivatives of oxanilide, benzalmalonates, α -cyanocinnamates and o-hydroxyphenylbenzotriazoles, having a reactive phenolic group(s). These additives are suggested for use with polymer coating materials.

[0006] U.S. Pat. No. 5,338,319 discloses a polyamide stabilization system containing a copper complex, and water soluble oxalic acid diaryl amide.

[0007] U.S. Patent No. 5,380,774 discloses a polyamide molding composition containing a hindered phenolic antioxidant, phosphite, hindered amine stabilizer and optionally a benzotriazole.

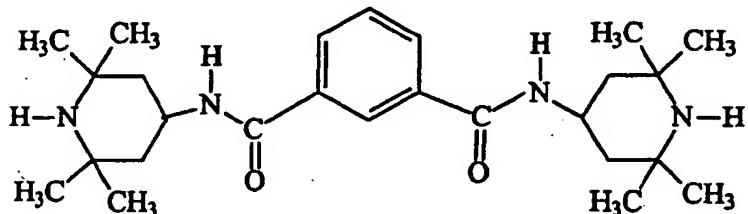
[0008] European Publication number EP 0379470 discloses compounds used to improve the dyeability of polyamide. The compounds include a compound containing a sterically hindered amine group, trimesic acid trialkylamide, pyromellitic acid tetra-alkylamide, triamino substituted triazine, a hydroxyl-reactive dyestuff. Compounds containing a sterically hindered amine group are preferably



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where R is hydrogen, C₁₋₈ alkyl, C₁₋₈ alkoxy, or COR₅.

- 40 [0009] WIPO publication WO 97/43335 discloses a stabilizer for polyamide having the structure:



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[0010] Polyamide polymers are characterized by high tensile strength, Young's modulus and abrasion resistance, however polyamides require stabilization against thermo-oxidative and photodegradation to sustain these properties in demanding use applications. Polyamides must accept lightfast dyeing systems in addition. Discoloration of the resin matrix is highly critical and may cause shade changes in colored articles, such as fibers and moldings of polyamide polymers. Therefore, further improvement in the stabilization systems for these materials is desirable in end uses such as fibers, and generally in high temperature processing of polyamide melts.

[0011] The polyamide structure contains labile hydrogens and exerts reductive or basic properties which are problematic from the standpoint of colorants. For example, in polyamide, organic red dyes undergo significant loss of shade upon thermal processing at the typical melt processing temperatures for polyamide. It would be desirable to minimize the tendency of polyamide to exert this influence on dyes while at the same time protecting the polymer matrix from loss of the desired physical properties.

[0012] The evolution of polyamide stabilization systems has advanced from earlier recommended hindered phenolic types, for example Irganox® 1098, to a combination of an antioxidant and a benzotriazole, such as the combination of Irganox® 1098 with Tinuvin® 234, well established in the art. Lightfast and photochemically stable dyed polyamide fibers containing oxalic acid di-aryl amides with a copper complex are known, for example from the aforementioned U.S. Pat. No. 5,338,319. These are applied in aqueous dye baths. In contrast, the present invention is directed to additives which are dispersed within the polyamide matrix by melt-processing at high temperatures above the glass transition, as in extrudates or moldings. WO 97/43335 provides a stabilizer which can be added in the melt to polyamide and improves heat stability, light stability, chemical stability, and dye affinity, however improvements are still sought.

[0013] Highly synergistic stabilization systems for polyamides have been found which exhibit unexpected further improvements for polyamide. These systems exhibit significant technical and economic advantages, even at low effective additive concentrations.

SUMMARY OF INVENTION

[0014] In accordance with the fundamental aspect of the invention there is provided a process for stabilizing polyamide polymers against the damage effected by light, heat and/or oxidation, even without the use of copper complexes, by using specified additives which act synergistically in polyamides. The process comprises incorporating the additive combinations selected from (I), (II) or (III), under melt processing conditions with polyamide, and forming a uniform and intimate mixture. The components, on a weight basis (I), (II) and (III) are:

(I) from 0.01% to 1% of a sterically hindered phenyl phosphonite (Ia) with from 0.05% to 5% of an oxanilide, (Ib), preferably non-hydroxy substituted, (together (Ia) + (Ib)), or (Ia) together with from 0.05 to 5% of an aromatic di- or tri- carbonyl (ester) amide compound, defined below containing at least one, and preferably two hindered amine moieties (Ic) in particular relationship with the carbonyl group, (together (Ia) + (Ic)). The weight ratio of (Ia) to (Ib) is from 1:3 to 1:10 and the weight ratio of (Ia) to (Ic) is from 1:3 to 1:10. The preferred weight ratios for both (Ia) to (Ib) as well as (Ia) to (Ic) is 1:3 to 1:6. The more preferred weight ratios of (Ia) to (Ib) and (Ia) to (Ic) are from 1:4 to 1:6 and most preferably 1:5;

(II) from 0.05 to 5% of an oxanilide (IIa), preferably non-hydroxy substituted, together with from 0.05 to 5% of an aromatic di- or tri- carbonyl (ester) amide compound containing at least one and preferably two hindered amine moieties (IIb), (together (IIa) + (IIb)). The weight ratio of (IIa) to (IIb) is from 1:5 to 5:1, preferably 1:3 to 3:1, and most preferably 1:1; and

(III) from 0.01% to 1% of a sterically hindered phenyl phosphonite, from 0.05 to 5% of an oxanilide, preferably non-hydroxy substituted, and from 0.05 to 5% of an aromatic di- or tri- carbonyl (ester) amide compound containing at least one, preferably two hindered amine moieties. The weight ratio of phosphonite to oxanilide is from 1:5 to 5:1, preferably 1:5 to 1:1, and most preferably 1:5 to 1:3

wherein the percentages represents grams of additives used per hundred grams of polymer.

[0015] The invention defines the basis, furthermore, for the effective and unexpected improved efficiency in the use of organic colorants with polyamide, such as solvent soluble, or polymer soluble organic dyes, for example, acid dispersed dyes, azo and diazo dyes, phthalocyanine blues and greens, diarylide yellows, perylenes, quinacridone, and the like. Therefore another aspect of the invention is a colored manufactured article such as a molding, extruded article or fiber of polyamide containing the stabilizer system (I), (II) or (III), in combination with one or more organic colorants.

DETAILED DESCRIPTION OF INVENTION

[0016] Polyamides stabilized according to the invention include the synthetic aliphatic, semi-aromatic and aromatic polyamides. Examples include nylon 6, nylon- 6.6, nylon-6.10, nylon 6.11, nylon-6.12, nylon 11, nylon 12, and copolymers such as nylon-6.6/6, nylon-6.6/6.10, 6/11, 6/12; polyether-polyamide block copolymers, poly(m-phenyleneisophthalamide), poly(p-phenyleneterephthalamide), and the like all of which are commercially available from a variety of sources. The method includes using the stabilization system with formulated polyamides, including conventional impact toughened polyamide, and reinforced polyamides containing glass fiber, mineral and glass/mineral combinations. The

invention can be practiced with polyamides regardless of the polymer morphology. Amorphous, semi-crystalline or highly crystalline polyamides as well as blends of different crystallinity are benefited. Commercially available polyamide resins are known from sources such as BASF, DSM, Evansville, Ind., E I duPont de Nemours, Wilmington, Del.; formulated polyamides are available, for example, from Ferro Corp., Cleveland, Ohio, Hanna Engineered Materials, Bethlehem, Pa., Hüls America, Somerset, N.J., and A. Schulman, Akron, Ohio; reinforced polyamides are available from DSM RIM Nylon, Westlake, Ohio.

[0017] The following amounts of each component selected according to the invention are critical to synergistic improvements and are indicated as weight percentage representing grams of additives used per hundred parts of the polyamide. The invention can be practiced by incorporating the additive system directly into heated zones of a compounding extruder containing the polyamide to be stabilized or by first forming an intermediate masterbatch compound which contains the selected system in a carrier polymer which is a polyamide or a polymer which is intimately compatible with or dispersible in polyamide, or by other methods commonly used in the art to incorporate additives into polymers.

[0018] According to the selection of ((Ia) + (Ib)) above, from 0.01 to 1% of sterically hindered phenyl phosphonite is used, preferably the usage level is from 0.05% to 0.5% with the most preferred level from 0.05 to 0.2% of the phosphonite. The specified oxanilide used in either (I) or (II) is possible at a level of from 0.05% to 5%, preferably at a level of from 0.2% to 2%, and most preferably at a level of from 0.3% to 1%.

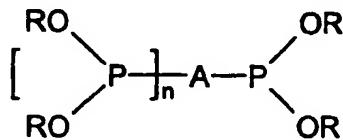
[0019] The aromatic di- or tri-carbonyl (ester)amide compound containing one or more piperidine groups is used generally in an amount of from 0.05% to 5%, preferably from 0.2% to 2% and most preferably from 0.3% to 1%.

[0020] According to (I) above the weight ratio of (Ia) to (Ib) or the ratio of (Ia) to (Ic) is 1:3 to 1:10, especially 1:4, 1:5, 1:6, 1:7, 1:8, and 1:9. According to ((IIa) + (IIb)) above, the weight ratio of (IIa) to (IIb) is from 1:5 to 5:1, especially 1:4, 1:3, 1:2, 1:1, 2:1, 3:1, and 4:1, with the most preferred weight ratio of from 1:2 to 2:1.

[0021] The sterically hindered phenyl phosphonites used herein have the following general structure:

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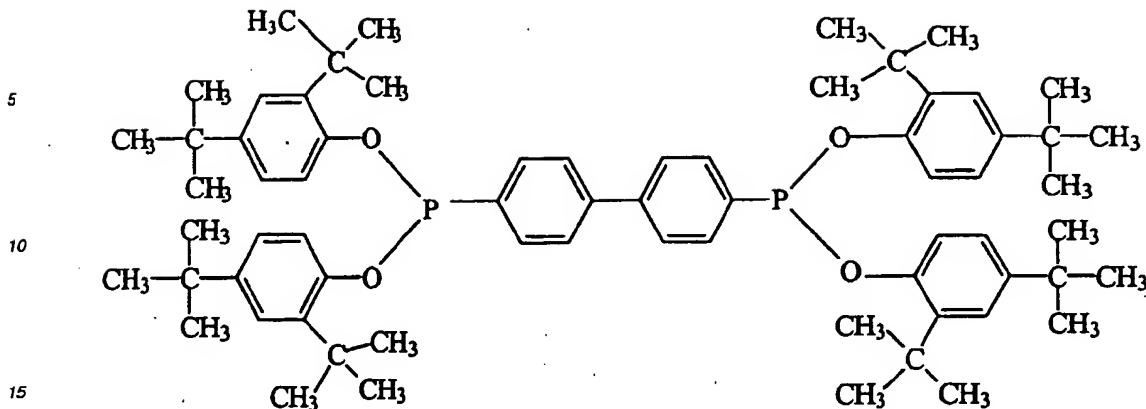
wherein each R is independently selected from unsubstituted phenyl or a substituted phenyl with one, two or three C₁₋₁₂alkyl groups, such as methyl, ethyl, propyl, butyl, isobutyl, tert.-butyl groups, linear or branched pentyl, hexyl, octyl or nonyl groups or cumyl moieties. Preferably R is a di-tert.-butyl phenyl group (most preferably at the 2,4-position), n is 0 or 1 and A is a predominantly aromatic mono- or difunctional C₅ to C₃₀ moiety optionally containing further heteroatoms, for example is a monofunctional or difunctional residue of a phenyl, diphenylether, diphenyl, diphenylmethane, or dibenzofuran group. Most preferably A is a diphenyl residue, n = 1 and R is a 2,4-di-tert-butyl phenyl group or 2,4-di-tert-butyl-5-methyl phenyl group. The term "sterically hindered" refers to the presence of at least one substituent different from hydrogen, in ortho position to the functional center of the respective group.

[0022] The most preferred phenyl phosphonites are the condensation products of sterically hindered alkyl phenols, such as 2,4-di-tertiary butyl phenol, with the Friedel-Crafts reaction product of diphenyl and PCl₃. An exemplary condensation product A-2 has the following structure:

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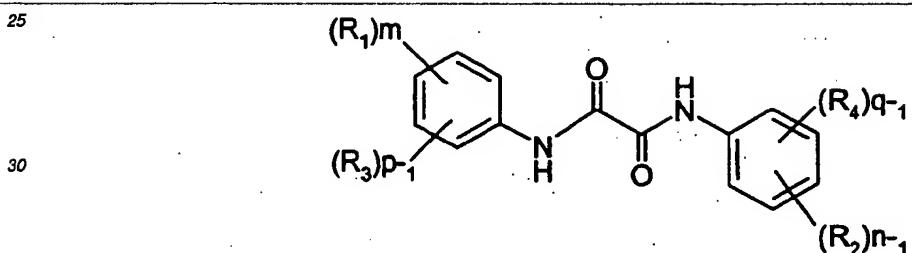
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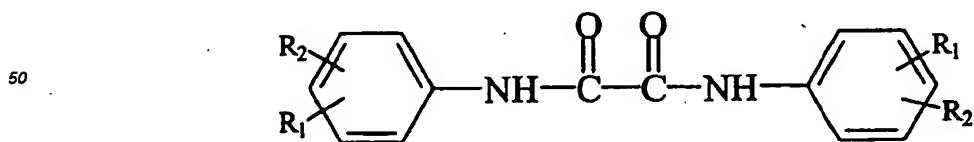
- [0023] The preferred sterically hindered phosphonite containing A-2 as main component is Sandostab® P-EPQ available from Clariant.

[0024] The oxalic acid di-aryl amides (oxanilides) used in the invention include, for example, compounds having the following structures:



- [0025] R₁ and R₂ are each independently of the other hydrogen, unsubstituted C₁-C₁₈alkoxy or C₁-C₁₈ alkoxy which is substituted by halogen, hydroxy, C₁-C₅ alkoxy, carboxyl groups, carbamyl groups or C₁-C₁₂ alkoxycarbonyl groups, or is C₃-C₅ alkenyloxy, unsubstituted benzyloxy or benzyloxy which is substituted by halogen or C₁-C₅alkyl, aliphatic acyloxy containing up to 18 carbon atoms, unsubstituted benzyloxy or benzoyloxy which is substituted by halogen or C₁-C₄alkyl, or a radical -A-SO₃M, wherein A is a direct bond or a divalent radical of formula -O-Q-, and Q is unsubstituted or hydroxy-substituted C₁-C₆alkylene, M is hydrogen or alkali metal, R₃ and R₄ are each independently of the other hydrogen, halogen, C₁-C₁₂alkyl, haloalkyl, phenyl or phenyl-C₁-C₅alkyl, or two radicals R₃ and/or R₄ in ortho-position each together form a fused 6-membered aromatic carbon ring, and wherein m and n are 1 or 2 and p and q are 1, 2 or 3.

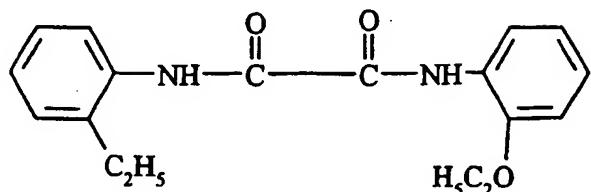
[0026] The preferred oxalic acid di-aryl amide is non-hydroxy substituted, for example:



- 55 wherein R₁ and R₂ are each independently, identical or different substituents selected from hydrogen, linear, branched or cyclic alkyl or alkoxy groups having from 2 to 12 carbon atoms and from 1 to 4 oxygen atoms; preferably one R₁ group on each ring is hydrogen, one R₂ group is ethoxy and the other is a C₂ to C₁₂ alkyl group, both located at the 5 position of each ring.

[0027] The most preferred oxanilides have such structures as:

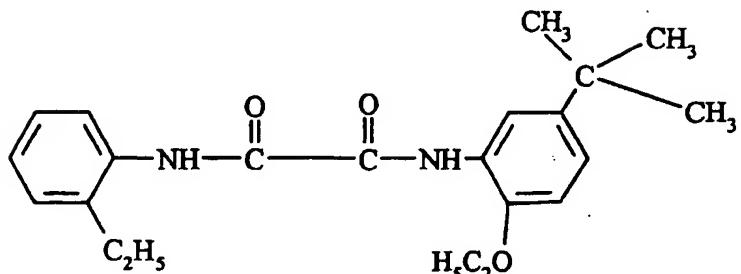
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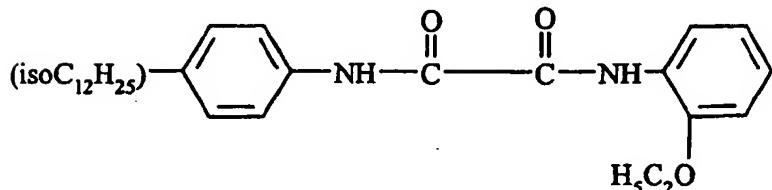
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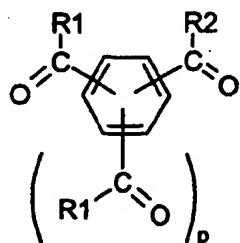


[0028] Commercially available non-hydroxyl substituted oxanilides include Sanduvor® VSU and 3206, and Tinuvin® 312 and 315, and the like. The most preferred oxanilide is commercially available from Clariant as Sanduvor® VSU.

[0029] The process according to (Ib) or (II) includes the use of an aromatic di- or tri-carbonyl compound which is an aromatic amide or aromatic ester-amide (aromatic (ester)amide) containing at least one hindered amine group adjacent to a carbonyl carbon. The structure is given by:

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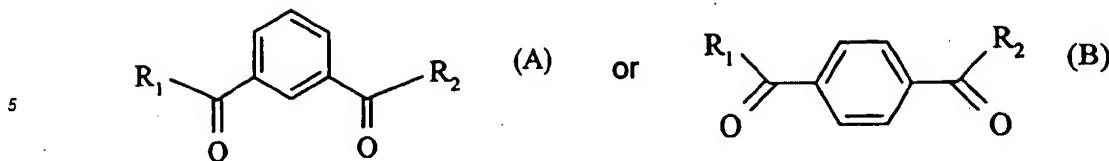
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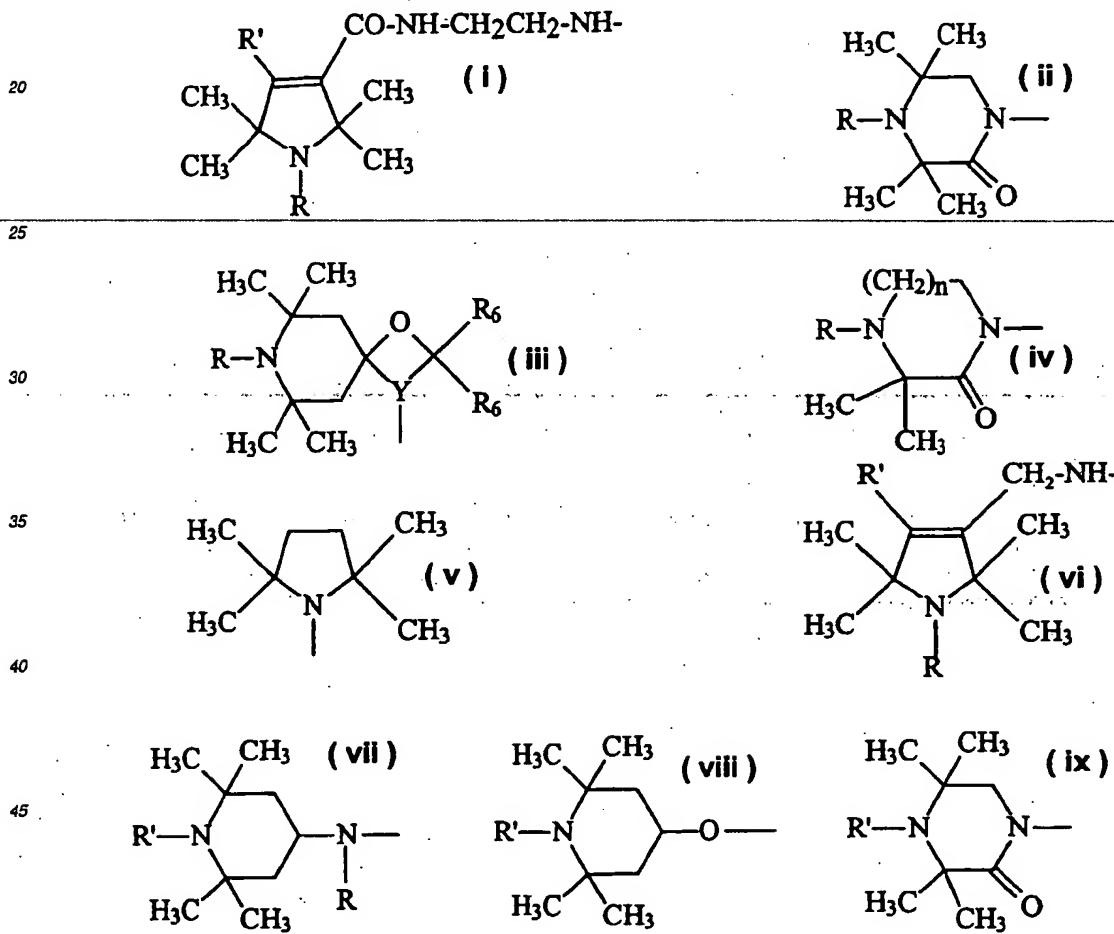
50 where p 0 or 1, and R¹ and R² is defined as for (A) and (B) below.

[0030] When p is 0, the aromatic di-carbonyl (ester)amide is (A) and (B).

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10 wherein at least one of R₁ and R₂ are independently, identical or different sterically hindered amino groups adjacent to the carbonyl group and the group(s) which is not a hindered amino group includes independently of the other, unsubstituted C₁-C₁₈alkoxy or alkamino which may optionally be substituted by halogen, hydroxy, C₁-C₅ alkoxy, carboxyl groups, carbamyl groups or C₁-C₁₂ alkoxy carbonyl groups, or is C₃-C₅ alkenyloxy, unsubstituted benzyloxy or benzylloxy which is substituted by halogen or C₁-C₅alkyl, aliphatic acyloxy containing up to 18 carbon atoms, unsubstituted benzyloxy or benzoyloxy which is substituted by halogen or C₁-C₄alkyl. Exemplary hindered amino groups include those such as (i) - (ix),

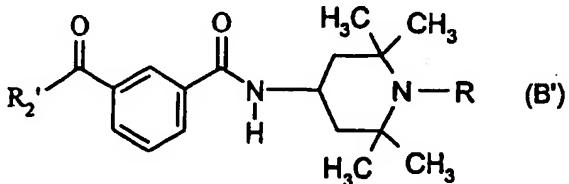


50 wherein R and R' are independently either hydrogen, C₁-C₁₂alkyl, C₁-C₈alkoxy, or -COR₃, where R₃ is hydrogen, C₁-C₆alkyl, phenyl, -COO(C₁-C₄alkyl) or NR₁₅R₁₆, where R₁₅ and R₁₆ are independently hydrogen, C₁-C₁₂alkyl, C₅-C₆cycloalkyl, phenyl or (C₁-C₁₂alkyl)phenyl; or R₁₅ and R₁₆ together with the N-atom to which they are attached form a five- to seven-membered ring which may contain an additional N- or O-atom (preferably forming a piperidine or morpholine ring); n is 0 or 1 (structure iv); Y is the group -NCO or -OCN, where CO forms part of the cyclic structure; each R₆ of structure (iii) independently is selected from hydrogen, C₁-C₁₂alkyl or phenyl provided only one R₆ can be phenyl, or both groups R₆ together form the group -(CH₂)_m; where m is 2 to 11, -C(CH₃)₂, -C(CH₃)₂-CH₂-CH₂- or -C(CH₃)CH₂CH₂CH(CH₃)-.

[0031] R₁ or R₂ groups other than hindered amino groups, include, independently C₁₋₂₀ alkyl, C₁₋₂₀ alkoxyalkyl, C₁₋₂₀ hydroxyalkyl, C₁₋₂₀ alkenyl, substituted C₁₋₂₀ alkenyl groups, C₁₋₂₀ alkoxy-C₁₋₂₀-alkyl groups, C₁₋₂₀-oxy-N-C₁₋₂₀-alkyl groups, C_{7-C₉} phenylalkyl unsubstituted or mono-, di- or tri-substituted on the phenyl by C_{1-C₄}alkyl; or aliphatic or aromatic C_{1-C₁₀}acyl.

- 5 [0032] A preferred aromatic dicarbonyl compound is (B'):

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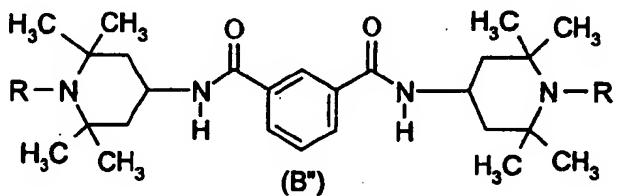


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wherein R_{2'} is independently C₁₋₂₀ alkyl, C₁₋₂₀ alkylamino, optionally further substituted C₁₋₂₀ alkoxyalkyl, C₁₋₂₀ hydroxyalkyl, C₁₋₂₀ alkenyl, substituted C₁₋₂₀ alkenyl groups, C₁₋₂₀-alkoxy-C₁₋₂₀-alkyl groups, C₁₋₂₀-oxy-N-C₁₋₂₀-alkyl groups, -N-cyclic-C₁₋₁₀-alkyl groups, and cyclic-N-C₁₋₁₀-alkyl groups -COR₄, where R₄ is hydrogen, C₁₋₆alkyl, phenyl, -C₁₋₂₀ COO(H or C₁₋₄alkyl).

- 20 [0033] More preferred is the aromatic dicarbonyl compound (B''):

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wherein each R in B' and B'' is independently either hydrogen, C_{1-C₁₈} alkyl, O, OH, CH₂CN, C_{1-C₁₆}alkoxy, alkoxy-alkylenoxyalkyl, C_{5-C₁₂}cycloalkoxy, C_{3-C₆}alkenyl, C_{7-C₉} phenylalkyl unsubstituted or mono-, di- or tri-substituted on the phenyl by C_{1-C₄}alkyl; or aliphatic or aromatic C_{1-C₁₀} acyl, or -COR₄, where R₄ is defined as above.

- 35 [0034] Examples of alkyl groups having not more than 20 carbon atoms are methyl, ethyl, propyl, isopropyl, butyl, 2-butyl, isobutyl, t-butyl, pentyl, 2-pentyl, hexyl, heptyl, octyl, 2-ethylhexyl, t-octyl, nonyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, hexadecyl and octadecyl; exemplary branched alkyl groups are isopropyl, isobutyl, t-butyl, 2,2-dimethylpropyl, 2-methylpropyl, cyclohexylmethyl, cyclohexylethyl, 2-ethylhexyl, 3,5,5-trimethylhexyl, 2-hexylundecyl, and 5,7,7-trimethyl-2-(1,3,3-trimethylbutyl)hexyl; exemplary cyclic alkyl groups are cyclohexyl.

40 [0035] Examples of alkoxy(alkyl) groups are methoxy, ethoxy, propoxy, isoproxy, butoxy, isobutoxy, pentoxy, isopen-toxy, hexoxy, 2-ethylhexyloxy, heptoxy, octoxy, decyloxy, dodecyloxy, tetradecyloxy, hexadecyloxy and octadecyloxy, nonadecyloxy. Preferred alkoxy examples are C_{6-C₁₂}alkoxy, in particular heptoxy and octoxy.

45 [0036] Examples of the alkoxyalkylenoxyalkyl groups are C₁₋₂₀-alkoxy-C₁₋₅-alkylenoxy-C₁₋₂₀-alkyl groups. The C₁₋₂₀-alkoxy groups and C₁₋₂₀-alkyl groups include those mentioned as examples of the alkoxyalkyl groups. Examples of the C₁₋₅-alkylenoxy groups include -CH₂O-, -CH₂CH₂O-, -CH₂CH₂CH₂O-,

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[0037] Particularly, C₁₋₁₂-alkoxy-C₁₋₅-alkylenoxy-C₁₋₁₂-alkyl groups are more preferred.

55 [0038] Examples of C_{5-C₁₂}cycloalkyl R₃ unsubstituted or mono-, di- or tri-substituted by C_{1-C₄}alkyl are cyclopentyl, methylcyclopentyl, dimethylcyclopentyl, cyclohexyl, methylcyclohexyl, dimethylcyclohexyl, trimethylcyclohexyl, t-butyl-cyclohexyl, cyclooctyl, cyclodecyl and cyclododecyl. Unsubstituted or substituted cyclohexyl is preferred.

[0039] Examples of C_{5-C₁₂}cycloalkoxy R₁ are cyclopentoxy, cyclohexoxy, cycloheptoxy, cyclooctoxy, cyclodecyloxy and cyclododecyloxy, including as applied to R₃. Cyclopentoxy and cyclohexoxy are preferred.

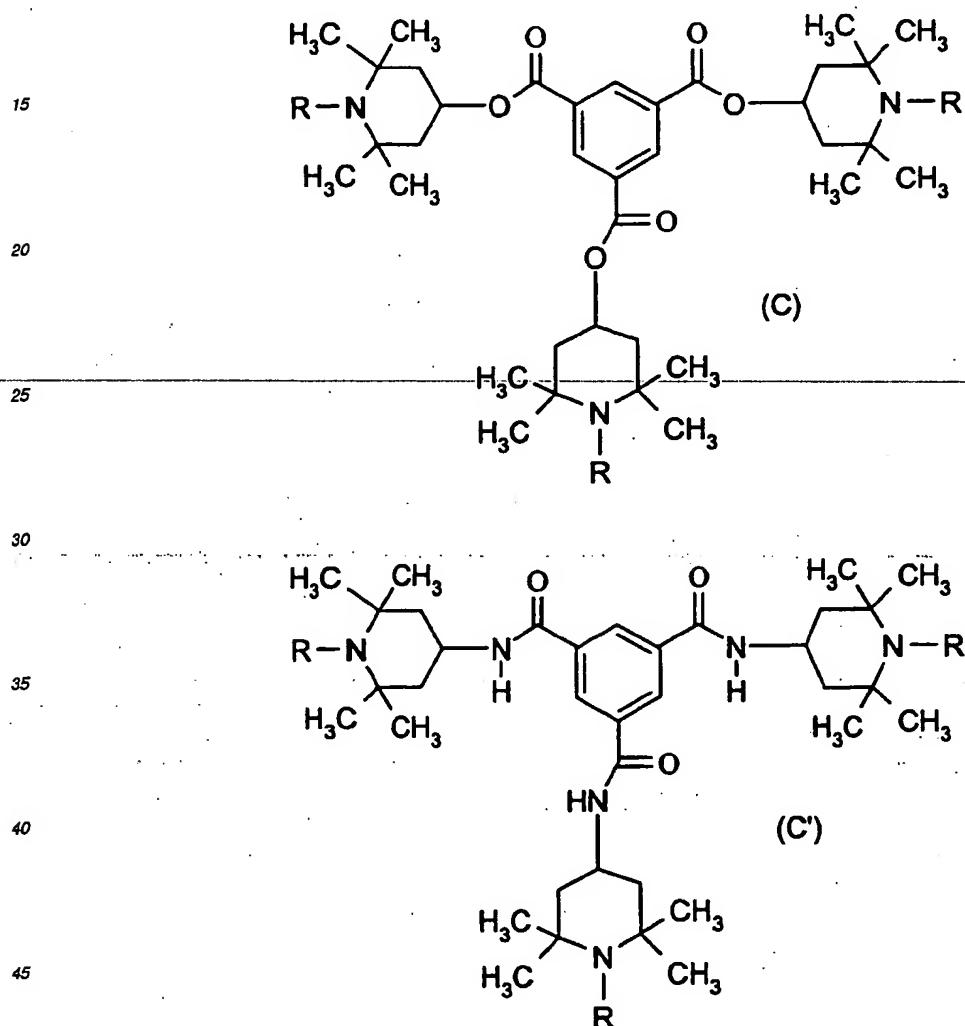
[0040] Examples of C₃-C₆ alkenyl are allyl, 2-methylallyl, butenyl and hexenyl, 3-butenyl, and 10-undecenyl; examples of branched alkenyl groups are 1-methyl-2-propenyl, 3-methyl-3-but enyl. Allyl is preferred.

[0041] Examples of C₇-C₉ phenylalkyl unsubstituted or mono-, di- or tri-substituted on the phenyl by C₁-C₄alkyl are benzyl, methylbenzyl, dimethylbenzyl, trimethylbenzyl, t-butylbenzyl and 2-phenylethyl. Benzyl is preferred.

5 [0042] Examples of aliphatic and aromatic C₁-C₁₀ acyl are C₁-C₈alkanoyl or C₃-C₈-alkenoyl, for example, acetyl, propionyl, butyryl, isobutyryl, pentanoyl, benzoyl, hexanoyl, heptanoyl, octanoyl, 2-ethylhexanoyl, acryloyl and crotonyl. Acetyl is preferred.

[0043] When more than one hindered amine group is present they can be identical or different, and preferably are identical hindered amine groups.

10 [0044] Exemplary aromatic tri-carbonyl compounds are the following C and C'.

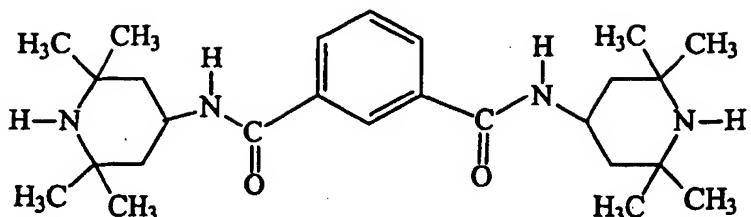


50 wherein R is defined as in B"

[0045] The most preferred aromatic (ester) amide is an aromatic di-amide having the following structure:

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and is commercially available as NYLOSTAB S-EED from Clariant.

[0046] The aromatic di- or tri-carbonyl (ester)amide with at least one hindered amino group can have remaining non-hindered amino groups R₁ and R₂ groups in structures (A) or (B) selected from radicals of substituted or unsubstituted linear or branched alkyl amine(s) or alkoxy groups having 8 to 20 carbon atoms, such as octyl amine and dodecylamine or the corresponding alcohols; substituted or unsubstituted aromatic amine, aminocarboxylic acid or corresponding lactam, also oligomers of about 3 to 20 repeating units of the aromatic dicarbonyl and a diamine or aminocarboxylic acid.

[0047] Conventional additives may optionally be present in the polyamide which is modified according to the invention, for example, primary thermal stabilizers, light stabilizers, lubricants, pigments, reinforcing and/or non-reinforcing fillers, and the like. Conventional primary antioxidants are suggested for inclusion herein at conventional amounts. Antioxidants include, for example alkylated monophenols, alkylated hydroquinones, alkylidenebisphenols, hindered alkyl benzyl compounds, acylaminophenols, esters of β -(3,5-di-tert-butyl-4-hydroxyphenyl) propionic acid, and the like which are disclosed in U.S. Pat. No. 5,051,459 incorporated herein by reference. A commercially available antioxidant is IRGANOX® 1098 from Ciba Specialty Chemicals.

[0048] UV-absorbers include, for example, the class of benzotriazoles such as 2-(2'-hydroxyphenyl)benzotriazoles, for example 2-(2'-hydroxy-5'-methylphenyl)benzotriazole, 2-(3',5'-di-tert-butyl-2'-hydroxyphenyl)benzotriazole, 2-(5'-tert-butyl-2'-hydroxyphenyl)benzotriazole, 2-[2'-hydroxy-5'-(1,1,3,3-tetramethylbutyl)phenyl]benzotriazole, 2-(3'-5'-di-tert-butyl-2'-hydroxyphenyl)-5-chlorobenzotriazole, 2-(3'tert-butyl-2'-hydroxy-5'-methyl-phenyl)-5-chlorobenzotriazole, 2-(3'-sec-butyl-5'-tert-butyl-2'-hydroxyphenyl)benzotriazole, 2-(2'-hydroxy-4'-octyloxy-phenyl)benzotriazole, 2-(3',5'-di-tert-amyl-2'-hydroxy-phenyl)-benzotriazole, 2-(3',5'-bis(9,9-dimethylbenzyl)-2'-hydroxyphenyl)benzotriazole; mixture of 2-(3'-tert-butyl-2'-hydroxy-5'-(2-octyloxycarbonylethyl)phenyl)-5-chlorobenzotriazole, 2-(3'-tert-butyl-5'-(2-ethyl-hexyloxy)carbonylethyl)-2'-hydroxyphenyl)-5-chlorobenzotriazole, 2-(3'-tert-butyl-2'-hydroxy-5'-(2-methoxycarbonylethyl)phenyl)-5-chlorobenzotriazole, 2-(3'-tert-butyl-2'-hydroxy-5'-(2-methoxycarbonylethyl)phenyl)benzotriazole, 2-(3'-tert-butyl-2'-hydroxy-5'-(2-ethyl-hexyloxy)carbonylethyl)-2'-hydroxyphenyl)-benzotriazole, 2-(3'-dodecyl-2'-hydroxy-5'-methylphenyl)-benzotriazole, and 2-(3'-tert-butyl-2'-hydroxy-5'-(2-isooctyloxycarbonylethyl)-phenylbenzotriazole, 2,2'-methylenebis[4-(1,1,3,3-tetramethylbutyl)-6-benzotriazol-2-ylphenol]; transesterification product of 2-[3'-tert-butyl-5'(2-methoxycarbonylethyl)-2'-hydroxyphenyl]benzo-triazole with polyethylene glycol 300; [R-CH₂CH₂-COO(CH₂)₃]₂ where R = 3'-tert-butyl-4'-hydroxy-5'-2H-benzotriazol-2-ylphenyl; [R-CH₂CH₂-COO(CH₂CH₂O)_n]₂ where n = 1 to 5 and R = 3'-tert-butyl-4'-hydroxy-5'-2H-benzotriazol-2-ylphenyl. A suitable commercially available benzotriazole is Norbloc® 7966, and 6000, from Jensen Pharma., Inc. Benzotriazoles are disclosed in U.S. Pats. 4,335,155, 4,405,749, and 4,528,311 which are incorporated herein by reference.

[0049] Further UV-absorbers include the class of ortho-hydroxyphenyl triazines such as 2-(2-hydroxyphenyl)-1,3,5-triazines, for example 2,4,6-tris(2-hydroxy-4-octyloxyphenyl)-1,3,5-triazine, 2-(2-hydroxy-4-octyloxyphenyl)-4,6-bis(2,4-dimethylphenyl)-1,3,5-triazine, 2-(2,4-dihydroxyphenyl)-4,6-bis-(2,4-dimethylphenyl)-1,3,5-triazine, 2-(2,4-dihydroxy-phenyl)-4,6-bis-(2,4-dimethylphenyl)-1,3,5-triazine, 2,4-bis(2-hydroxy-4-propyloxyphenyl)-6-(2,4-di-methylphenyl)-1,3,5-triazine, 2-(2-hydroxy-4-octyloxyphenyl)-4,6-bis(4-methyl-phenyl)-1,3,5-triazine, 2-(2-hydroxy-4-dodecycloxyphenyl)-4,6-bis(2,4-dimethylphenyl)-1,3,5-triazine, 2-[2-hydroxy-4-(2-hydroxy-3-butyloxypropyloxy)phenyl]-4,6-bis(2,4-di-methylphenyl)1,3,5-triazine and 2-hydroxy-4-(2-hydroxy-3-octyloxypropyloxy)-phenyl]-4,6-bis(2,4-dimethylphenyl)-1,3,5-triazine. A suitable commercially available triazine is CYASORB® 1164, from Cytec Corp.

[0050] Hindered benzoate UV-absorbers include commercially available materials such as CYASORB® 2908 from Cytec, and FERRO® AM 340 from Ferro Corp.

[0051] Further, in this specification, where a range is given, the figures defining the range are included therein.

[0052] Further, any group capable of being linear, branched or cyclic is linear, branched or cyclic.

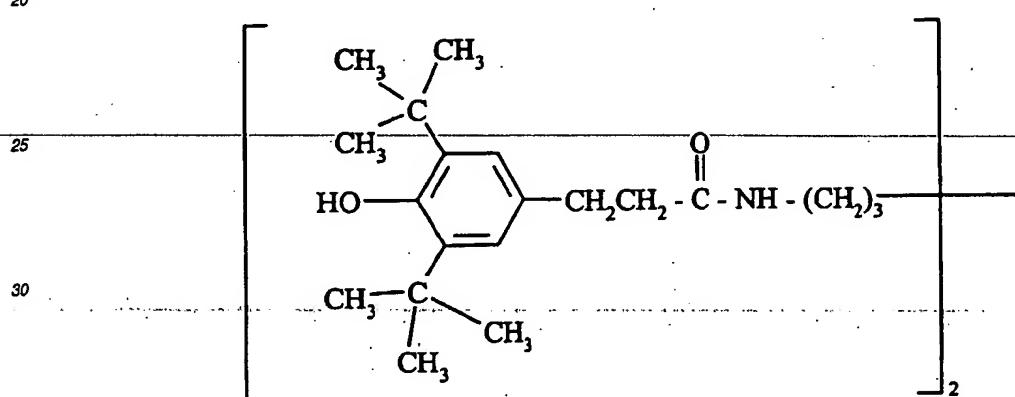
[0053] For the avoidance of doubt, in this specification tert-butyl means tertiary butyl or $\text{C}(\text{CH}_3)_3$.

[0054] The following examples illustrate the invention in its various aspects. All parts and percentages are expressed by weight, representing grams of additives used per hundred grams of polyamide.

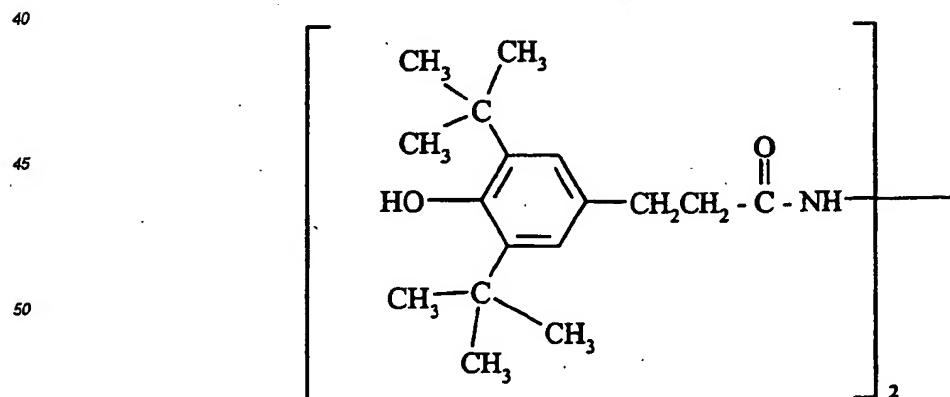
EXAMPLES

[0055] In the comparative examples and in the exemplary non-limiting embodiments of the invention described hereunder, the following types of polyamide-based polymers and additives are used:

- 5 1) PA-1 A polyamide 6 of the low-viscosity type suitable especially for injection molding and commercially available as Capron[®] 8202 NL from Allied Signal Corp.
- 10 2) PA-2 A transparent, amorphous, polyamide-polyether block copolymer suitable for injection molding and commercially available from EMS-CHEMIE, Domat/Ems, Switzerland.
- 15 3) PA-3 A semi-loss polyamide 6 containing 0.3 wt.% of TiO₂ commercially available as Nylon 6 SD 1130 from DSM.
- 20 4) PS-1 A sterically hindered phosphonite processing stabilizer commercially available as Sandostab[®] P-EPQ from Clariant, Muttenz, Switzerland.
- 25 5) PS-2 A primary hindered phenolic antioxidant commercially available as Irganox[®] 1098 from Ciba Specialty Chemicals, Basle, Switzerland and which has the following structure:



- 30 6) PS-3 An antioxidant which is a compound of the following formula:

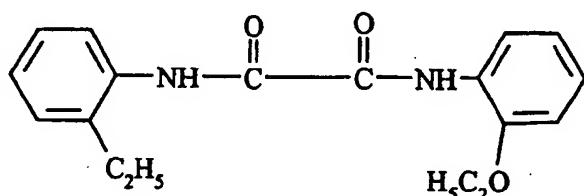


35 40 45 50 55 PS-3 is commercially available as Irganox[®] MD 1024 from Ciba Specialty Chemicals, Basle, Switzerland.

- 7) LS-1 A non-hydroxyl substituted oxanilide UV-absorber commercially available as Sanduvor® VSU from Clariant, having the following structure:

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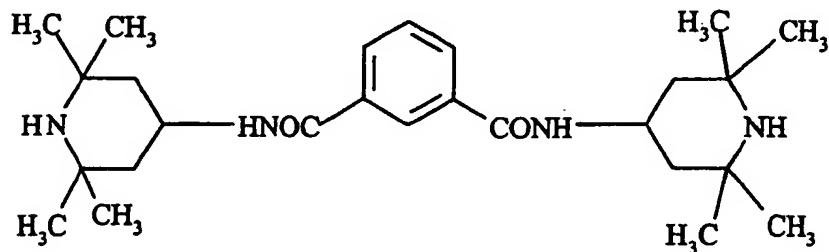


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- 8) LS-2 An aromatic hindered piperidine UV-stabilizer commercially available as Nylostab® S-EED from Clariant, and which is a compound of the following formula:

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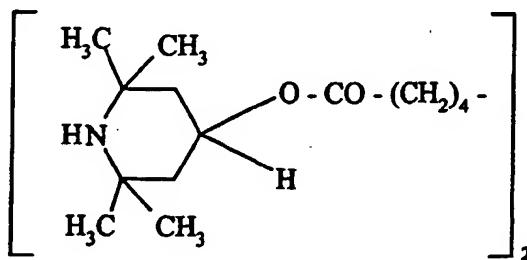
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- 9) LS-3 A UV-stabilizer which is commercially available as Tinuvin (RTM) 770 from Ciba Specialty Chemicals in Basle, Switzerland and which is a compound of the following formula:

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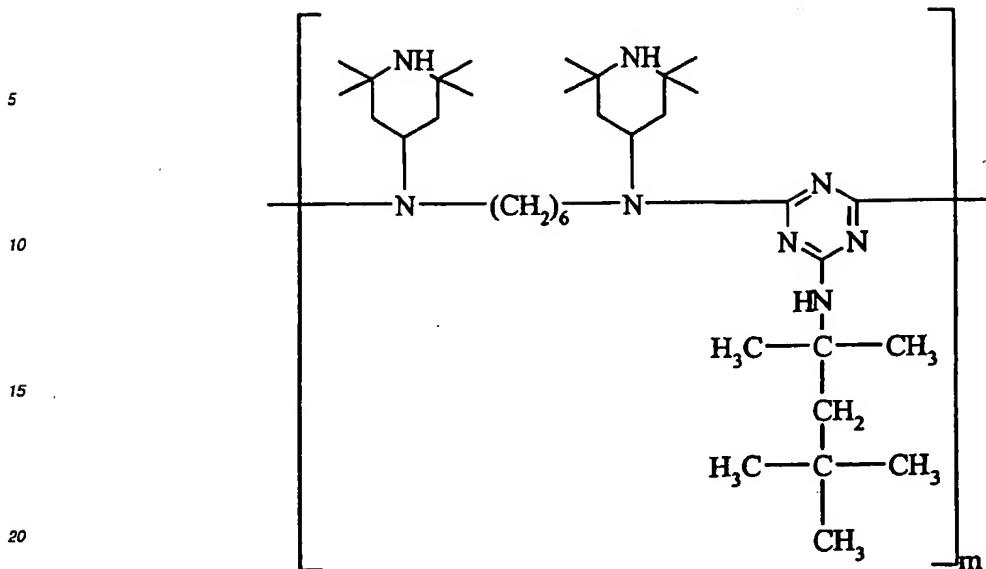
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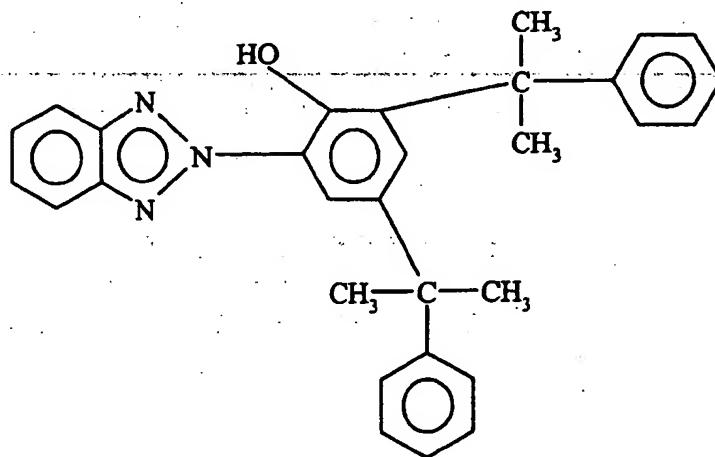


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- 10) LS-4 A UV-stabilizer which is commercially available as Chimassorb (RTM) 944 from Ciba Specialty Chemicals in Basle, Switzerland or as Sanduvor (RTM) 3944 from Clariant in Muttenz, Switzerland and which is a compound of the following formula:

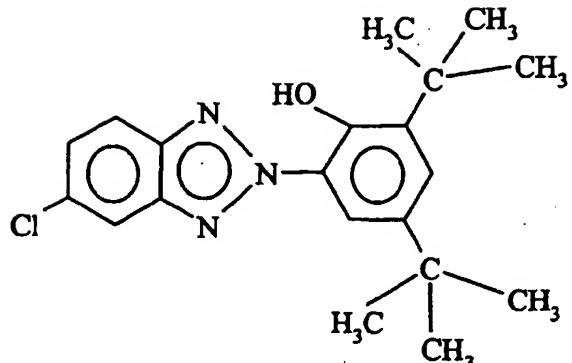


- 11) LS-5 A UV-absorber which is commercially available as TINUVIN® 234 from Ciba Specialty Chemicals in Basle, Switzerland and which is a compound of the following formula:



- 12) LS-6 A UV-absorber which is commercially available as TINUVIN® 327 from Ciba Specialty Chemicals, Basle, Switzerland and which is a compound of the following formula:

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- [0056] The tested samples were prepared as follows. According to the formulations described hereunder, the different types of polyamides were first melt compounded with the tested additives and extruded at temperatures of from 250-280° Celsius, followed by pelletizing into homogeneous pellets. These pellets were formed into panels 1 mm thick by injection molding at temperatures of from 280 to 300° Celsius, followed by either punching of the tested samples to 35 mm x 40 mm or by direct injection molding of tensile bar specimens.
- [0057] The color determination is reported in terms of the Yellowness Index (YI) determined in accordance with the ASTM E-313 Yellowness Test. The higher the value, the lower the color quality, i.e. the stronger the discoloration.
- [0058] In addition, the color is measured according to the regulations of the CIE (International Commission on Illumination) and in accordance with the ISO 7724/3 and DIN 6174 standards. The higher the positive value of b*, the more pronounced the yellowing.
- [0059] The tensile strength is measured in accordance with the ASTM D-638 standard.
- [0060] The accelerated weathering test (Atlas WOM 65 WRC) is carried out according to the DIN 53387-A standard.
- [0061] The results of the tests carried out with stabilizer compositions according to the invention and with stabilizers according to the prior art (for comparative purposes) applied to different types of polyamides are summarized in the tables 1-3.

35

TABLE 1

Yellowness index and b*-value of the various formulations containing PA-1 as polymer after processing into the tested samples.			
No.	Formulation	Yellowness index	b*-value
1	Ref. Sample	1.36	0.61
2	0.1 % PS-1	0.71	0.31
3	0.5 % LS-1	4.03	1.79
4	0.1 % / 0.5% of PS-1 / LS-1	-2.42	-1.06
5	0.5 % LS-2	3.87	1.70
6	0.1% / 0.5% of PS-1 / LS-2	-4.57	-1.98
7	0.5 % LS-5	17.20	7.86
8	0.1 % / 0.5% of PS-1 / LS-5	15.07	6.85
9	0.5 % LS-6	14.29	6.52
10	0.1% / 0.5% of PS-1 /LS-6	12.96	5.88

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- [0062] Test numbers 4 and 6 of table 1 are formulations according to the invention; test numbers 7-10 of table 1 are comparative tests.

[0063] The effect of the processing stabilizer PS-1 to improve the values of yellowing can be seen from the results shown in table 1. From the data of this table it can also be gathered that the preferable use of the UV-stabilizers LS-1 and LS-2 is clearly appropriate, although up to now some minor worsening had to be accepted in the preparation of light-stabilized polyamides.

5 [0064] Therefore, the synergism observed with the stabilizer composition according to the invention is totally unexpected. In spite of the high thermal stress during the processing of the polyamides into the tested samples, yellowing does not occur at all. The measured values of the Yellowness Index and of the b*-value do not only achieve the level of the reference sample but exceed the quality of the reference sample in a significant manner.

10 [0065] The comparative tests show that the use of the processing stabilizer PS-1 in combination with other light stabilizers than those according to the invention, such as LS-5 and LS-6 lead only to minor improvements of the yellowing values which are generally increased versus the reference as soon as a light stabilizer is present in the matrix to be stabilized. The synergism caused by the stabilizer composition according to the invention is not observed in other cases, i.e. the synergism is limited to the composition according to the invention only.

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TABLE 2

Tensile strength (TS) and b*-value of various formulations containing PA-2 as polymer after exposure to accelerated weathering (WOM 65 WRC) for 1800 hours.			
No.	Formulation, always with 0.2% PS-2	b*-value	TS (% of the original value)
1	Reference sample	7.85	29
2	0.6 % LS-1	4.04	84
3	0.6 % LS-2	4.71	39
4	0.6 % LS-3	4.65	49
5	0.6 % LS-4	5.40	40
6	0.3% / 0.3% of LS-1 / LS-2	2.62	96
7	0.3 % / 0.3 % of LS-1 / LS-3	3.05	88
8	0.3 % / 0.3 % of LS-1 / LS-4	3.58	82

35 [0066] Test number 6 of table 2 is a formulation according to the invention; test numbers 7 and 8 of table 2 are comparative tests.

40 [0067] If light stabilizers are not used the weathering causes a markedly increase of the yellowing value as well as a distinct worsening of the mechanical properties of PA-2 (cf: reference sample). Further, the data of table 2 shows that the high efficacy of the UV-stabilizer LS-1 exceeding clearly the effectiveness of the UV-stabilizers LS-2, LS-3 and LS-4 in view of the yellowing value b* and the tensile strength. As it is obvious from table 2, the combination of LS-1 and LS-2 surprisingly results in a pronounced synergism enabling the production of polyamides having a markedly improved light fastness, i.e. resistance against yellowing, as well as a better UV-fastness, i.e. retention of the mechanical properties.

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TABLE 3

Table 3 illustrates b*-values and condition of the tested samples containing PA-3 as polyamide after exposure to accelerated weathering (WOM 65 WRC) at the time indicated.			
	Formulation always with 0.25 wt. % PS-2	b*- value	Time of embrittlement
1	reference sample	n.d. ¹	3000 h
2	0.5% / 0.25% LS-1 / LS-2	2.6	> 4000 h
3	0.5% / 0.25% LS-1 / LS-3	n.d. ¹	3000 h
4	0.5% / 0.25% LS-1 / LS-4	3.3	> 4000 h
5	0.5% / 0.25% LS-5 / LS-2	5.6	> 4000 h
6	0.5% / 0.25% LS-5 / LS-3	4.6	> 4000 h
7	0.5% / 0.25% LS-5 / LS-4	4.7	> 4000 h

n.d.¹sample decomposed- not measurable

[0068] Test number 2 of Table 3 is a formulation according to the invention; test numbers 3 to 7 of Table 3 are comparative tests.

[0069] In this test series, samples were exposed to accelerated weathering for up to 4000 hours followed by measurement of b*-yellowing, and evaluation of brittleness. The stabilized polyamide containing LS-1 with LS-2 according to the invention showed outstanding efficiency with regard to yellowing resistance and retention of mechanical properties. The stabilizer combination No. 2 of LS-1/LS-2 far out-performed those combinations of LS-1/LS-3 and LS1/LS-4 and accordingly any combinations of LS-5 evidencing an unexpected and unique result.

Claims

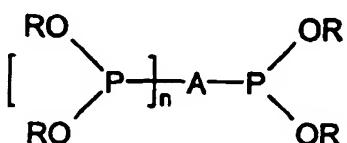
1. A process for stabilizing polyamide polymers against the damage effected by light, heat and/or oxidation comprising incorporating therein by melt-processing a synergistic additive system resulting in a uniform and intimate mixture with said polyamide, said additive system is selected from the groups consisting of (I) or (II), wherein

(I) comprises from 0.01% to 1% by weight of a sterically hindered phenyl phosphonite (Ia), and from 0.05% to 5% by weight of an oxanilide (Ib), or said (Ia) together with from 0.05% to 5% by weight of an aromatic di- or tri-carbonyl compound containing at least one hindered amine moiety adjacent to said carbonyl(s) (Ic); and

(II)-comprises from 0.05% to 5% by weight of said substituted oxanilide (IIa), and from 0.05% to 5% by weight of said aromatic di- or tri-carbonyl compound containing at least one hindered amine moiety adjacent to said carbonyl(s) (IIb),

wherein the percentages represent grams of additives used per hundred grams of polymer.

2. The process according to claim 1 wherein the weight ratio of (Ia) to (Ib) is from 1:3 to 1:10, the weight ratio of (Ia) to (Ic) is from 1:3 to 1:10, and the weight ratio of (IIa) to (IIb) is from 1:5 to 5:1.
3. The process according to claim 1 wherein said sterically hindered phenyl phosphonite has the following structure:



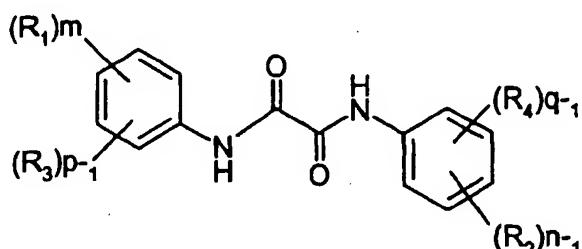
wherein each R is independently unsubstituted phenyl or a substituted phenyl with one, two or three alkyl groups

each having from 1 to 12 carbon atoms or a substituted phenyl with one or two cumyl groups, n is 0 or 1, and A is a predominantly aromatic, mono- or difunctional C₅-C₃₀ moiety optionally containing further heteroatoms.

4. The process according to claim 1 wherein said oxanilide has the following structure:

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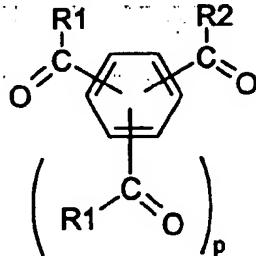
R₁ and R₂ are each independently of the other hydrogen, unsubstituted C₁-C₁₈alkoxy or C₁-C₁₈ alkoxy which is substituted by halogen, hydroxy, C₁-C₅ alkoxy, carboxyl groups, carbamyl groups or C₁-C₁₂ alkoxy carbonyl groups, or is C₃-C₅ alkenyloxy, unsubstituted benzyloxy or benzyloxy which is substituted by halogen or C₁-C₅ alkyl, aliphatic acyloxy containing up to 18 carbon atoms, unsubstituted benzoyloxy or benzoyloxy which is substituted by halogen or C₁-C₄ alkyl, or a radical of formula -A-SO₃M, wherein A is a direct bond or a divalent radical of formula -O-Q-, and Q is unsubstituted or hydroxy-substituted C₁-C₆alkylene, M is hydrogen or alkali metal, R₃ and R₄ are each independently of the other hydrogen, halogen, C₁-C₁₂alkyl, haloalkyl, phenyl or phenyl-C₁-C₅alkyl, or two radicals R₃ and/or R₄ in ortho-position each together form a fused 6-membered aromatic carbon ring, and wherein m and n are 1 or 2 and p and q are 1, 2 or 3.

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5. The process according to claim 1 wherein (II) is selected and wherein said aromatic di- or tri-carbonyl (ester)amide contains at least one hindered amine group adjacent to the carbonyl carbon, said aromatic di- or tri-carbonyl (ester)amide is given by:

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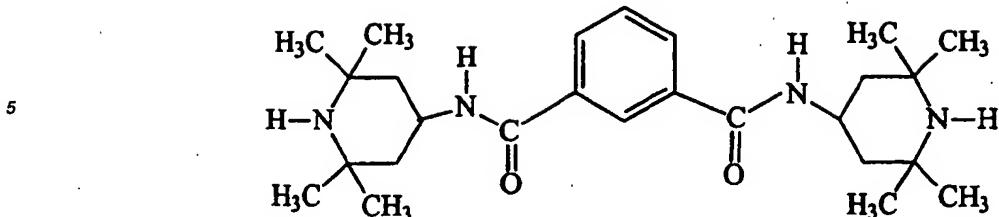


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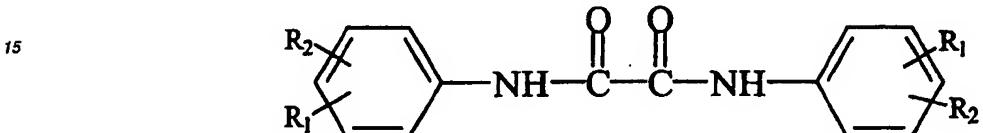
where p is 0 or 1, wherein at least one of R₁ and R₂ are independently, identical or different sterically hindered amino groups and the group(s) R₁ and R₂ other than sterically hindered amino group include, independently of the other, unsubstituted C₁-C₁₈alkoxy or alkamino which may optionally be substituted by halogen, hydroxy, C₁-C₅ alkoxy, carboxyl groups, carbamyl groups or C₁-C₁₂ alkoxy carbonyl groups, or is C₃-C₅ alkenyloxy, unsubstituted benzyloxy or benzyloxy which is substituted by halogen or C₁-C₅ alkyl, aliphatic acyloxy containing up to 18 carbon atoms, unsubstituted benzoyloxy or benzoyloxy which is substituted by halogen or C₁-C₄ alkyl.

- 50 6. The process according to claim 1 wherein said (ester amide) has the structure:

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and said oxanilide has the following structure:



20 wherein R₁ and R₂ are each independently, identical or different moieties selected from hydrogen, linear, branched or cyclic alkyl or alkoxy groups having from 2 to 12 carbon atoms and from 1 to 4 oxygen atoms.

- 25
7. The process according to claim 1 wherein said additive system is selected from the group (III) which is from 0.01% to 1% by weight of a sterically hindered phenyl phosphonite, from 0.05% to 5% by weight of said oxanilide and from 0.05% to 5% by weight of said aromatic di- or tri-carbonyl compound, and wherein the percentages represent grams of additives used per hundred grams of polymer.
 - 30 8. A polyamide article selected from the group consisting of molded articles, extruded articles and fibers which comprise polyamide, optionally an organic dye, and in intimate homogeneous mixture therewith, an additive system selected from the group consisting of (I), (II) and (III) as defined in claim 1 or 7.
 9. The article according to claim 8 comprising an additive system as defined in any one of the preceding claims 1 to 7.

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EUROPEAN SEARCH REPORT

Application Number
EP 98 11 7816

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
X	DE 42 33 973 A (SANDOZ AG) 15 April 1993 * abstract * * page 2, line 30 - line 52 * * page 3, line 49 - page 4, line 56 * * page 5, line 10 - line 11 * * examples * * claims 1,3,6,7 *	1	C08K5/00 C08K5/20 //(C08K5/00, 5:20,5:3435, 5:5333)
A	US 5 147 935 A (VOLMER SOPHIE ET AL) 15 September 1992 * abstract * * examples * * claims *	1-9	
A	EP 0 430 650 A (HOECHST CELANESE CORP) 5 June 1991 * abstract * * claims *	1-9	
A	DE 195 37 140 A (SANDOZ AG) 11 April 1996 * abstract * * page 5, line 25 - line 40 * * examples * * claims *	1,9	TECHNICAL FIELDS SEARCHED (Int.Cl.6) C08K
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	22 January 1999	Pollio, M	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons A : technological background B : non-written disclosure P : intermediate document & : member of the same patent family, corresponding document	
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**ANNEX TO THE EUROPEAN SEARCH REPORT
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EP 98 11 7816

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22-01-1999

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
DE 4233973	A	15-04-1993		FR 2682392 A GB 2260764 A IT 1263005 B JP 5214254 A		16-04-1993 28-04-1993 23-07-1996 24-08-1993
US 5147935	A	15-09-1992		BE 1003675 A AT 110401 T DE 69103547 D DE 69103547 T EP 0443663 A JP 5239343 A		19-05-1992 15-09-1994 29-09-1994 13-04-1995 28-08-1991 17-09-1993
EP 0430650	A	05-06-1991		US 5380774 A AT 134207 T CA 2027802 A DE 69025379 D DE 69025379 T JP 3181561 A MX 174069 B		10-01-1995 15-02-1996 29-05-1991 28-03-1996 04-07-1996 07-08-1991 19-04-1994
DE 19537140	A	11-04-1996		ES 2116896 A FR 2725451 A GB 2293827 A,B IT RM950655 A JP 8113672 A		16-07-1998 12-04-1996 10-04-1996 09-04-1996 07-05-1996

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